Continuous Filament Glass Fibre and Human Health

SUMMARY

Continuous filament glass fibres (CFGF) produced by GlassFibreEurope member companies have diameters greater than 6 microns. These fibres therefore have filament diameters above the respirable size of 3 microns or less, thus minimising the potential for any chronic pulmonary effects associated with exposure to fibres.

Customers can confirm the diameter of the fibre that they purchase from their supplier. The irritation that can possibly be caused by these fibres is the result of mechanical abrasion, which can be minimised by good industrial hygiene practices.

Manufacturers and their customers should continue to use approved safety and health practices to ensure safe use of our products. Work practices and procedures should be in place to minimise dust generation. Local exhaust ventilation should be used if necessary to minimise and/or keep airborne dust levels below recommended limits. A government approved dust respirator should be used if airborne concentrations exceed regulatory and recommended limits, if irritation occurs, or if the workers choose to do so for personal comfort. Exposure assessments should be conducted, as appropriate, to ensure exposures are within recommended limits.

1. CHARACTERISTICS

1.1. Introduction

Glass fibres have been manufactured and placed on the market for more than 60 years. During this time, they have become one of the world’s most useful and beneficial man-made materials.

While they have numerous uses and applications, glass fibres are generally produced in two basic forms: wool-type fibres, referred to most commonly as glass wool or glass fibre insulation, and CFGF, produced in long, continuous strands or filaments.

1.2. Continuous Filament Glass Fibre (CFGF) Products and Applications

CFGF products are produced and supplied in a variety of forms: roving, chopped strands, yarns, mats, fabrics, tissues, milled fibres etc. The main end-use is the reinforcement of thermosetting and thermoplastic resins. These composite materials are used in a wide variety of applications.

The leading markets for composite materials are the automotive and transport sectors, the electrical/electronics industry, and the building industry. Other markets include pipes and tanks, agricultural equipment, industrial machinery, wind-turbine blades, and the sports, leisure and marine sectors. Another important end-use is the
manufacture of textiles that are used in similar markets to composites, though clearly for different applications. The main market for glass textiles in the electronics industry is in the production of printed wiring boards.

1.3. Man-Made (Synthetic) Vitreous Fibres (MMVF / SVF)
Glass fibres are categorised within a group of man-made materials historically referred to as man-made mineral fibres (MMMF). However, a more appropriate name is man-made vitreous fibres (MMVFs) or synthetic vitreous fibres (SVFs), reflecting the glassy, non-crystalline nature of the material. The glass used to produce the fibres is made by melting sand and other inorganic materials under highly controlled conditions.

1.4. Composition of CFGF
The predominant glass composition for continuous filament glass fibre is known as E-glass. E-glass is a member of the family calcium-aluminium-silicate glasses.

Boron is generally a major element of E-glass, with sodium and potassium maintained at low levels to give acceptable electrical properties. In recent years, however, alternative E-glass formulations, without boron and fluorine, have been developed for use in most industrial applications, except for printed wiring boards or aerospace applications.

For some applications requiring specific properties, such as higher mechanical strength, higher temperature resistance, improved resistance to corrosion, resistance to alkali in cement, or high dielectric properties, other glass families like C, D, R, AR and S are also used to manufacture CFGF.

1.5. Manufacturing CFGF
Glass fibres are a high technology product. CFGF is produced by a continuous drawing process through calibrated holes or bushings at constant speed, thus leading to a very narrow variation in filament diameter.

The diameter of the filaments does not differ significantly from the mean or nominal diameter. The standard deviation of the mean diameter of CFGF products is typically less than 10% of the nominal diameter. The typical diameter of CFGF products manufactured by GlassFibreEurope member companies ranges from greater than 6 to 25 microns with the majority of the products being 9 microns or larger in diameter.

The manufacturing process also provides a parallel orientation to the continuous filaments constituting the fibre bundles.

Further processing of CFGF products does neither generate a change in diameter, nor in the parallel orientation of filament bundles.

CFGF are not considered nano-materials as defined by the European Commission (A natural, incidental or manufactured material containing particles, in an unbound state or as an aggregate or as an agglomerate and where, for 50% or more of the particles in the number size distribution, one or more external dimensions is in the size range 1 nm - 100 nm.)
2. HEALTH AND SAFETY ASPECTS OF CFGF

2.1. Inhalation

Airborne dust of CFGF can be inhaled. However, the potential for inhaled glass fibres to cause any health hazard depends on its “respirability”, i.e., its potential to enter the lower regions of the lung. Indeed, the essential feature of a health and safety assessment for the product is to determine whether it is possible for the product to cause lung disease through respiration.

According to the WHO definition, respirable fibres have a diameter (d) smaller than 3 microns, a length (l) larger than 5 microns and an l/d-ratio greater than or equal to 3. Fibres with diameters greater than 3 microns, which is the case for continuous filament glass fibre, do not reach the lower respiratory tract and, therefore have no possibility of causing serious pulmonary disease.

CFGF do not possess cleavage planes which would allow them to split length-wise into fibres with smaller diameters, rather they break across the fibre, resulting in fibres which are of the same diameter as the original fibre with a shorter length and a small amount of dust. However in the specific case of the milled fibre product, the amount of dust can be more significant and specific respiratory protection equipment (dust mask) is recommended.

Microscopic examination of dust from highly chopped and pulverised glass showed the presence of small amounts of respirable dust particles. Among these respirable particles, some were fibre-like in terms of l/d ratio (so-called “shards”). Under the microscope, however, it can be clearly observed that they are not regularly shaped fibres but irregularly shaped particles with fibre-like dimensions. To the best of our knowledge, the exposure levels of these fibre-like dust particles measured in the GlassFibreEurope member companies’ manufacturing plants are in an order of magnitude of 50 to 1000 times below existing applicable limits.

2.2. Irritation

In contact with skin or eyes CFGF dust can cause a purely mechanical irritation (itching). This is definitely not an allergic reaction.

When sufficient amounts of CFGF are released into the air during manufacture and handling, some workers may experience temporary upper respiratory tract irritation. Like skin irritation, upper respiratory irritation is a mechanical reaction to the fibres. It is not an allergic reaction and the irritation generally does not persist. Such exposures to high concentrations of airborne fibres may result in temporary coughing and/or sneezing. These effects will subside after the worker is removed from exposure, and should have no further impact on his or her health or well-being.

By respecting the manufacturers’ safe use and handling instructions, these mechanical effects can readily be avoided.

As a general rule, the mechanical irritation caused by glass fibres disappears when the person ceases to be exposed to the product.
2.3. Human Epidemiology Studies
An important method for assessing the effects of a substance on humans is through epidemiological studies. Such studies typically examine large groups of people who have been exposed to the substance being studied.

Two major studies involving 21500 workers in the USA and Europe, conducted respectively by the University of Pittsburgh, School of Public Health, and the International Agency for Research on Cancer (IARC), showed no increase in lung cancer or non-malignant respiratory disease among persons working in glass fibre production. A smaller study was conducted among workers in a CFGF manufacturing facility in Canada with the same results.

Three epidemiological studies have been published on cohorts of people working in MMVF factories. The first one in Europe by Boffeta & al. (1997) on different types of MMVF concluded for two plants in Northern Ireland and Italy that there was no significant increase of different types of cancer compared to reference cohorts. The two other studies by Chiazze al. (1997) were specifically made in one plant producing CFGF in the USA. Chiazze concluded that there was no evidence of excess of cancer in the populations working in this plant for a long time (more than 15 years). References are mentioned below.

2.4. Classification and Regulatory Aspects
Several major reviews have been undertaken by various international expert organizations on the health and safety aspects of glass fibres. The first of these was conducted by the International Agency for Research on Cancer (IARC 1987). The purpose of the IARC review was to determine whether these fibres are carcinogenic to humans. At that time, IARC concluded that continuous filament glass fibres are not classifiable as to their carcinogenicity to humans (IARC classification Group 3). In October 2001, after a comprehensive review of more recent human epidemiology and animal toxicity data, IARC concluded that the classification of CFGF in Group 3 is appropriate, confirming that there is currently no evidence for the carcinogenicity of continuous filament glass fibres to humans (IARC 2002).

IARC groups man-made vitreous fibres (MMVF) into categories based on raw materials, production process, and end use. IARC noted, in its 2001 reclassification of MMVFs, that an additional category had been added to group those durable glass fibres produced by flame attenuation for special applications. IARC retained the Group 2B classification for what IARC termed “Special Purpose Fibres.” IARC gave as examples of these SPF: E-and 475 respirable glass fibres. IARC retained the Group 3 classification for continuous filament fibres, regardless of chemical composition. Continuous filament fibres differ from Special Purpose Fibres in their method of manufacture and end use. They may also have different composition. Thus, continuous filament E-glass fibres should not be confused with SPF E-glass fibres and are still classified as Group 3.

Environment Canada also completed a review of the scientific data for glass fibres. The purpose of the review was to assess both the hazards of glass fibres and the risk to humans and the environment presented by those fibres. It concluded for continuous filament glass fibres:

“Based principally on the likelihood that few respirable fibres are generated in the production and use of continuous filament and that concentrations in the general environment should be extremely small, it has been concluded that continuous glass filament is not entering the environment in quantities or under conditions that may constitute a danger in Canada to human life or health” (Environment Canada 1993).
The American Conference of Governmental Industrial Hygienists (ACGIH) has classified continuous filament glass fibres as not classifiable as human carcinogen. The ACGIH has established a TLV (Threshold Limit Value or recommended exposure limit) for glass fibre of 1 fibre per cubic centimetre of air for respirable fibres and 5 mg per cubic meter of air for inhalable glass fibre dust. These levels were established to prevent mechanical irritation of the upper airways. NTP (US National Toxicology Program) and OSHA (US Occupational Safety and Health Administration) do not list continuous filament glass fibres as a carcinogen.

According to the German Technical Rules for Hazardous Substances TRGS 905 (list of CMR substances), only WHO fibres (diameter less than 3 µm, length greater than 5 µm and a length/diameter ratio of greater than 3) are considered (suspected) carcinogens.

CGGF as produced by GlassFibreEurope member companies do not fulfil these criteria. All GlassFibreEurope produced continuous filament glass fibre types have a diameter larger than 3 µm and hence are not to be considered as (suspected) carcinogens under the definitions of the German TRGS. The KI-index given in the TRGS 905 does not apply to continuous filament glass fibre as produced by the GlassFibreEurope member companies.

Continuous filament glass fibres are not considered as a dangerous substance following the rules of the European CLP regulation and its subsequent amendments. This has been confirmed by the 1st ATP 790/2009 of the CLP regulation on Man-made Mineral Fibre where continuous filament glass fibres are not to be labelled either for toxicity, carcinogenicity, or irritation. Labelling is only applicable to glass or rock wool in certain circumstances and refractory ceramic fibres.

2.5. Industry Recommended Work Practices
While CFGF are safe to manufacture and handle, a number of general work practices should nevertheless be followed by those who are involved in these operations. In addition to preventive measures aiming to reduce the possibilities of generating dust or broken filaments, a series of protective measures in areas of high exposure are recommended: gloves, long-sleeved cloths, long-legged trousers, and dust masks especially for workers involved in cutting operations, cleaning or discharging of containers. It is furthermore recommended to measure, as appropriate, the number of fibres in the air to prevent high exposure levels to fibre or dust, in order to ensure compliance with existing exposure limits.

Years of airborne fibre sampling at GlassFibreEurope manufacturing facilities confirm that very low concentrations of respirable fibres may be present, but the concentrations are well below current recommended exposure limits. GlassFibreEurope will continue to conduct exposure monitoring to ensure proper work practices, engineering controls, and personal protective equipment (PPE) are in place to eliminate or minimise exposure risk.

GlassFibreEurope product information will continue to be reviewed and updated as needed, based upon the evaluation of work by different laboratories studying these subjects and the ongoing analysis of our products.
3. References


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